## **AMENDMENTS TO THE CLAIMS**

1. (original) A method for the synthesis of severely sterically hindered secondary aminoether alcohols of the formula

wherein R<sup>1</sup> and R<sup>2</sup> are each selected from the group consisting of alkyl, hydroxylalkyl radicals having 1 to 4 carbon atoms or in combination with the carbon atom to which they are attached they form a cycloalkyl group having 3 to 8 carbon atoms, and R<sup>3</sup> is selected from the group consisting of hydrogen, alkyl hydroxyalkyl radicals having 1 to 4 carbon atoms, and mixtures thereof, and R<sup>4</sup>, R<sup>5</sup>, R<sup>6</sup>, R<sup>7</sup>, R<sup>8</sup>, R<sup>9</sup>, R<sup>10</sup> and R<sup>11</sup> are the same or different and are selected from the group consisting of hydrogen, alkyl and hydroxyalkyl radicals having 1 to 4 carbons provided that at least one of R<sup>4</sup> or R<sup>5</sup> bonded to the carbon atom directly bonded to the nitrogen atom is an alkyl or hydroxyalkyl radical when R<sup>3</sup> is hydrogen, the process involving reacting an organic carboxylic acid or salt of a carboxylic acid of the formula

$$R^{12} - C - OY$$

wherein  $R^{12}$  is selected from the group consisting of alkyl radicals having 1 to 4 carbon atoms, aryl radicals bearing hydrogen or one or more  $C_1$ - $C_{10}$  alkyl groups substituted thereon, and mixtures thereof, and Y is selected from the group consisting of hydrogen, alkali metal, ammonium, and mixtures thereof, with a sulfonyl halide, a sulfuryl halide, a mixed sulfuryl ester halide, or a mixed sulfuryl amide halide of the formula

$$R^{14}SO_2X$$
,  $SO_2X_2$ ,  $R^{14}OSO_2X$ , or  $R_2^{14,14}NSO_2X$ 

wherein X is selected from the group consisting of F, Cl, Br, I, and mixtures thereof, and  $R^{14}$  and  $R^{14'}$  are the same or different and each is selected from the group consisting of alkyl radicals having 1 to 4 carbon atoms, haloalkyl radicals of the formula  $C_nH_{(2n+1)-w}Z_w$ 

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wherein n is 1 to 4, Z is selected from the group consisting of F, Cl, Br, I, and mixtures thereof, and w ranges from 1 to 5, and aryl radicals

$$R^{16}$$
 $R^{15}$ 
 $R^{17}$ 
 $R^{18}$ 
 $R^{19}$ 

wherein R<sup>15</sup>, R<sup>16</sup>, R<sup>17</sup>, R<sup>18</sup>, and R<sup>19</sup> are the same or different and are selected from hydrogen and alkyl radicals having 1 to 20 carbon atoms, and mixtures thereof, to yield acyl sulfonate material of the general formula

$$R^{12} = C - O - SO_{2}R^{14}$$

$$R^{12} = C - O - SO_{2} - O - C - R^{14}$$

$$R^{12} = C - O - SO_{2} - O - C - R^{14}$$

$$R^{12} = C - O - SO_{2} - O - R^{14}$$

$$R^{12} = C - O - SO_{2} - O - R^{14}$$

which is then reacted with a dioxane of the formula

$$\begin{array}{c}
R^{11} & O \\
R^{10} & O \\
R^{9} & R^{6}
\end{array}$$

wherein R<sup>4</sup>, R<sup>5</sup>, R<sup>6</sup>, R<sup>7</sup>, R<sup>8</sup>, R<sup>9</sup>, R<sup>10</sup>, and R<sup>11</sup> are the same or different and are selected from hydrogen, alkyl and hydroxyalkyl radicals having 1 to 4 carbons to yield

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or

which is then aminated with an alkylamine of the formula

$$R^{1}$$
 $H_{2}N - C - R^{2}$ 
 $R^{3}$ 

wherein R1, R2, and R3 are as previously defined to yield

which is then hydrolyzed with base to yield

2. (original) The method of claim 1 for the synthesis of severely sterically hindered secondary aminoether alcohols using sulfonyl halide of the formula  $R^{14}SO_2X$ .

- 3. (original) The method of claim 1 for the synthesis of severely sterically hindered secondary aminoether alcohols using sulfuryl halide of the formula  $SO_2X_2$ .
- 4. (original) The method of claim 1 for the synthesis of severely sterically hindered secondary aminoether alcohols using the mixed sulfuryl ester halide of the formula R<sup>14</sup>OSO<sub>2</sub>X.
- 5. (original) The method of claim 1 for the synthesis of severely sterically hindered secondary aminoether alcohols using the mixed sulfuryl amide halide of the formula  $R_2^{14,14}NSO_2X$ .
- 6. (currently amended) The method of according to any one of the preceding claims claim 1, 2, 3, 4 or 5 wherein R<sup>1</sup>, R<sup>2</sup> and R<sup>3</sup> are methyl radicals.
- 7. (currently amended) The method of according to any one of the preceding claims claim 1, 2, 3, 4 or 5 wherein R<sup>4</sup>, R<sup>5</sup>, R<sup>6</sup>, R<sup>7</sup>, R<sup>8</sup>, R<sup>9</sup>, R<sup>10</sup>, and R<sup>11</sup> are hydrogen.
- 8. (currently amended) The method of according to any one of the preceding claims claim 1, 2, 3, 4 or 5 wherein R<sup>15</sup>, R<sup>16</sup>, R<sup>18</sup>, and R<sup>19</sup> are hydrogen and R<sup>17</sup> is hydrogen or methyl.
- 9. (currently amended) The method of according to any one of the preceding claims claim 1, 2, 3, 4 or 5 wherein the base is selected from alkali metal hydroxide, alkali metal alkoxide, or alkali metal carbonate.
- 10. (currently amended) The method of according to any one of the preceding claims claim 1, 2, 3, 4 or 5 wherein Y is hydrogen or sodium.
- 11. (currently amended) The method of according to any one of the preceding claims claim 1, 2, 3, 4 or 5 wherein R<sup>1</sup>, R<sup>2</sup> and R<sup>3</sup> are methyl, R<sup>4</sup>, R<sup>5</sup>, R<sup>6</sup>, R<sup>7</sup>, R<sup>8</sup>, R<sup>9</sup>, R<sup>10</sup>, and R<sup>11</sup> are hydrogen, R<sup>15</sup>, R<sup>16</sup>, R<sup>18</sup>, and R<sup>19</sup> are hydrogen, R<sup>17</sup> is hydrogen or methyl, and Y is hydrogen, sodium, or ammonium.

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12. (currently amended) The method of according to any one of the preceding claims claim 1, 2, 3, 4 or 5 wherein the acyl sulfonate is made by reacting organic carboxylic acid or the salt of a carboxylic acid with the sulfonyl halide, sulfuryl halide, mixed sulfuryl ester halide or mixed sulfuryl amide halide at a temperature in the range of between about -20 to 200°C at a pressure between about 1 bar and 100 bars, the acyl sulfonate is reacted with the dioxane at a molar ratio of dioxane to acyl sulfonate in the range of 1:1 to 10:1 at a temperature of between about 50°C to about 200°C to yield a cleavage product, the cleavage product and the alkyl amine reacted at an amine to sulfonate group ratio ranging from about stoichiometric to about 10:1 at pressure of from about atmospheric (1 bar) to about 100 bars at temperature of from about 40°C to about 200°C, and the resulting aminated product is hydrolyzed with base at a temperature from about 20°C to about 110°C.

13. (currently amended) The method of according to any one of the preceding claims claim 1, 2, 3, 4 or 5 wherein the organic carboxylic acid or the salt thereof, the sulfonyl halide, sulfuryl halide, mixed sulfuryl ester halide or mixed sulfuryl amide halide and the dioxane care combined in a single step to produce a reaction mixture, the reaction mixture being heated at a temperature of between about 50°C to about 200°C to produce the cleavage product, the cleavage product and the alkylamine are reacted at am amine to cleavage product ratio ranging from about stoichiometric to about 10:1 at a pressure from about atmospheric (1 bar) to about 100 bars at a temperature of from about 40°C to about 200°C, the resulting aminated product being reacted with base at a temperature from about 20°C to about 110°C.

Applicants request that the present application be examined on the basis of the above presented amended claims.

Respectfully submitted,

Joseph J. Allocca

Attorney for Applicants Registration No. 27,766

Telephone Number: (908) 730-3629 Facsimile Number: (908) 730-3649

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ExxonMobil Research and Engineering Company P. O. Box 900 Annandale, New Jersey 08801-0900

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